

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Original) A peptide, comprising the sequence $S_1 - S_2 - S_3 - S_4 - S_5$, wherein S_1 comprises an amino acid chain from one to about four neutral or charged L- or D-configuration amino acid residues or a linear or branched alkyl, aryl, alkene, alkenyl or aralkyl chain; S_2 is absent or is a natural or unnatural aliphatic amino acid residue; S_3 is L- or D-Phe, Phe(4-Cl), Phe(2,4-diCl), Phe(3,4-diCl), Phe(4-NO₂), Phe(4-Me), Phe(4-Phenyl), Hphe, Pgl, Trp, Nal 1, Nal 2, Bip, Dip, Bpa, Ser(Bzl), Lys(Z), Lys(Z-2'Br), Lys(Bz), Thr(Bzl), Cys(Bzl), Tyr(BzlCl₂) or any natural or unnatural L- or D-amino acid with an aromatic side chain group, wherein the aromatic ring is optionally functionalized with halogen, alkyl or aryl groups; S_4 is L- or D-Lys, Arg, Orn, Dpr, Dbu, p-amino-Phe or any natural or unnatural amino acid with a positively charged side chain; S_5 comprises an L- or D-amino acid with an aromatic side chain.
2. (Original) The peptide of claim 1 wherein S_1 potentiates the intrinsic activity of the remainder of the peptide by providing an auxiliary or secondary receptor contact.
3. (Original) The peptide of claim 1 wherein S_1 comprises an acetyl group.
4. (Original) The peptide of claim 1 wherein S_2 is Gly or L- or D-Ala, Val, Leu or Nle.
5. (Original) The peptide of claim 1 wherein S_4 is an L-configuration cationic amino acid.
6. (Currently amended) The peptide of claim 1 wherein S_5 ~~the L- or D-amino acid with an aromatic side chain~~ is an L- or D-isomer of Phe, Phe(4-Cl), Phe(2,4-diCl), Phe(3,4-diCl), Phe(4-NO₂), Phe(4-Me), Phe(4-Phenyl), Hphe, Pgl, Trp, Nal 1, Nal 2, Bip, Dip, Bpa, Ser(Bzl),

Lys(Z), Lys(Z-2'Br), Lys(Bz), Thr(Bzl), Cys(Bzl), Tyr(BzlCl₂), an N-alkylated or arylated derivative of any of the foregoing, or a des-carboxyl amino acid corresponding to any of the foregoing.

7. (Original) The peptide of claim 1 wherein S₅ comprises one or more additional amino acids.

8. (Original) The peptide of claim 1 wherein S₅ comprises a terminal group.

9. (Currently amended) A peptide, ~~comprising~~ consisting of the sequence S₁ – S₂ – D-Phe(4-Cl) – S₄ – S₅, wherein

S₁ is heptanoyl, 2'-naphthylacetyl, 7'-amino-heptanoyl, 2'-chlorophenylacetyl, 3'-

chlorophenylacetyl, 4'-chlorophenylacetyl, 4'-phenylbutylaminocarbonyl, 3'-

phenylbutylaminocarbonyl, 4'-bromophenyl-acetyl, 3,4-dichlorophenyl-acetyl, 2,4-

dichlorophenyl-acetyl, 4-biphenyl-acetyl, 2-naphthoyl, Ph-(CH₂)₂NH, 3'-

phenylpropanecarbonyl, 2'-naphthoyl-Pip, 2'-naphthylacetyl, 2'-bromophenyl-acetyl, 4'-

CF₃phenyl-acetyl, 3'-CF₃phenyl-acetyl, 2'-CF₃phenyl-acetyl, 3',5'-CF₃phenylacetyl, 2',5'-

CF₃phenylacetyl, 4'-Mephenyl-acetyl, 3'-Mephenyl-acetyl, 2'-Mephenyl-acetyl, 7'-

aminoheptonoyl, beta-Ala, 4-aminoButyl, 5-aminoValeryl, 6-aminoCaproyl,

aminoTranexamyl, Cmpi or 3',4'-Cl₂phenylacetyl;

S₂ is absent or is Ser(Bzl), Ala, D-Ala, beta-Ala, Val, Leu, Chg, Aib, Tle, 1-amino-

1cyclohexanecarbonyl, Inp, CO(CH₂)₂NH, CO(CH₂)₂CO, Pip, MeThr(Bzl), Thr(Bzl) or D-

Thr(Bzl);

S₄ is Arg, D-Arg, (Nlys)Gly, Trp, Lys, homoLys, Dpr(beta-Ala), alpha-(N-amidino-4'-piperidine)Gly,

(4'-guanidino)Gly, (4'-guanidino)Phe, D-(4'-guanidino)Phe, beta-(N-amidino-4'-

peperidine)Ala or homo-Ala-4'-pip(N-amidino); and

S₅ is Trp, Trp-OH, Trp-NH₂, Trp-Cys-NH₂, D-Trp, D-Trp-NH₂, Trp-Val-NH₂, 3'-Pya-NH₂, Phe-NH₂,

MeTrp-NH₂, beta-Ala-Trp-NH₂, aminobutylamide, Nal 1-NH₂, D-Nal 1-NH₂, Nal 2-NH₂, D-

Nal 2-NH₂, Tic-NH₂, D-Tic-NH₂, 1'-aminoindan, 1'-aminoindane-1-carboxyl-NH₂, Aic-NH₂,

Atc-NH₂, Disc-NH₂, Tpi-NH₂, D-Tpi-NH₂, Tiq-NH₂, D-Tiq-NH₂, tryptamide, NMe-

tryptamide, alpha-Me-tryptamide, 2'-(4"-methylphenyl)ethylamide, 3',4'-Cl₂)phenylmethylamide, 3'-phenylpropylamide, 2',4'-dichlorobenzylamide, 3'-(1H-imidazol)propylamide, 4-phenyl-piperidine-4-carbonamide, 3-phenyl-1-propylamide, 2,4-dichlorophenethylamide, S-(-)-1-(2-naphthyl)ethylamide, S-(-)-1-(1-naphthyl)ethylamide, 2'-methylbenzylamide, 4'-methylbenzylamide, 2',2'-diphenylethylamide, 1-(2-pyridyl)piperazine, N-benzylmethylamide, histamide, R-(+)-1-(2-Naphthyl)ethylamide, Trp-Asp-NH₂, Trp-Asp-Phe-NH₂, Asp-Trp-NH₂, Ala-Trp-NH₂, Trp-Ala-NH₂, phenethylamide or Trp-Asp-OH.

10. (Original) The peptide of claim 9 consisting of the sequence:

7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Ala-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-Phe-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-NH₂,
 heptanoyl-Thr(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-MeTrp-NH₂,
 heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-MeTrp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-NMe-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-alpha-Me-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-S-(-)-1-(1-Naphthyl)ethylamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 1-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-D-Nal 2-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 2-NH₂,
 2'-naphthylacetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂,
 4'phenylbutyryl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂,
 3',4'-dichlorophenyl-acetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂, or
 3'-CF₃phenyl-acetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂.

11. (Original) A peptide, ~~comprising~~ consisting of the sequence 7'-amino-heptanoyl – S₂ – D-Phe(4-Cl) – S₄ – S₅, wherein

S₂ is absent or is Ser(Bzl), Ala, D-Ala, beta-Ala, Val, Leu, Chg, Aib, Tle, 1-amino-1cyclohexanecarbonyl, Inp, CO(CH₂)₂NH, CO(CH₂)₂CO, Pip, MeThr(Bzl), Thr(Bzl) or D-Thr(Bzl);

S₄ is Arg, D-Arg, (Nlys)Gly, Trp, Lys, homoLys, Dpr(beta-Ala), alpha-(N-amidino-4'-piperidine)Gly, (4'-guanidino)Gly, (4'-guanidino)Phe, D-(4'-guanidino)Phe, beta-(N-amidino-4'-peperidine)Ala or homo-Ala-4'-pip(N-amidino); and

S₅ is Trp, Trp-OH, Trp-NH₂, Trp-Cys-NH₂, D-Trp, D-Trp-NH₂, Trp-Val-NH₂, 3'-Pya-NH₂, Phe-NH₂, MeTrp-NH₂, beta-Ala-Trp-NH₂, aminobutylamide, Nal 1-NH₂, D-Nal 1-NH₂, Nal 2-NH₂, D-Nal 2-NH₂, Tic-NH₂, D-Tic-NH₂, 1'-aminoindan, 1'-aminoindane-1-carboxyl-NH₂, Aic-NH₂, Atc-NH₂, Disc-NH₂, Tpi-NH₂, D-Tpi-NH₂, Tiq-NH₂, D-Tiq-NH₂, tryptamide, NMe-tryptamide, alpha-Me-tryptamide, 2'-(4"-methylphenyl)ethylamide, 3',4'-Cl₂)phenylmethylamide, 3'-phenylpropylamide, 2',4'-dichlorobenzylamide, 3'-(1H-imidazol)propylamide, 4-phenyl-piperidine-4-carbonamide, 3-phenyl-1-propylamide, 2,4-dichlorophenethylamide, S-(-)-1-(2-naphthyl)ethylamide, S-(-)-1-(1-naphthyl)ethylamide, 2'-methylbenzylamide, 4'-methylbenzylamide, 2',2'-diphenylethylamide, 1-(2-pyridyl)piperazine, N-benzylmethylamide, histamide, R-(+)-1-(2-Naphthyl)ethylamide, Trp-Asp-NH₂, Trp-Asp-Phe-NH₂, Asp-Trp-NH₂, Ala-Trp-NH₂, Trp-Ala-NH₂, phenethylamide or Trp-Asp-OH.

12. (Original) The peptide of claim 11 consisting of the sequence

7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Ala-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-Phe-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-MeTrp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-NMe-Tryptamide,

7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-alpha-Me-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-S-(-)-1-(1-Naphthyl)ethylamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 1-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-D-Nal 2-NH₂, or
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 2-NH₂.

13. (Original) A peptide, ~~comprising~~ consisting of the sequence S₁ – S₂ – S₃ – S₄ – S₅, wherein

S₁ is heptanoyl, 2'-naphthylacetyl, 7'-amino-heptanoyl, 2'-chlorophenylacetyl, 3'-chlorophenylacetyl, 4'-chlorophenylacetyl, 4'-phenylbutylaminocarbonyl, 3'-phenylbutylaminocarbonyl, 4'-bromophenyl-acetyl, 3,4-dichlorophenyl-acetyl, 2,4-dichlorophenyl-acetyl, 4-biphenyl-acetyl, 2-naphthoyl, Ph-(CH₂)₂NH, 3'-phenylpropanecarbonyl, 2'-naphthoyl-Pip, 2'-naphthylacetyl, 2'-bromophenyl-acetyl, 4'-CF₃phenyl-acetyl, 3'-CF₃phenyl-acetyl, 2'-CF₃phenyl-acetyl, 3',5'-CF₃phenylacetyl, 2',5'-CF₃phenylacetyl, 4'-Mephenyl-acetyl, 3'-Mephenyl-acetyl, 2'-Mephenyl-acetyl, 7'-aminoheptonoyl, beta-Ala, 4-aminoBytyl, 5-aminoValeryl, 6-aminoCaproyl, aminoTranexamyl, Cmpi or 3'4'-Cl₂phenylacetyl;

S₂ is absent or is Ser(Bzl), Ala, D-Ala, beta-Ala, Val, Leu, Chg, Aib, Tle, 1-amino-1cyclohexanecarbonyl, Inp, CO(CH₂)₂NH, CO(CH₂)₂CO, Pip, MeThr(Bzl), Thr(Bzl) or D-Thr(Bzl);

S₃ is Phe, D-Phe, Phe(4-Cl), D-Phe(4-Cl), Phe(3-Cl), D-Phe(3-Cl), Phe(2-Cl), D-Phe(2-Cl), D-Phe(3,4-diCl), MePhe, D-MePhe, D-Tic, D-Tpi, D-Nal 2, Arg, D-Phe(3,4-F₂), D-Tiq, D-Me(homo)Phe or D-EtPhe;

S₄ is Arg, D-Arg, (Nlys)Gly, Trp, Lys, homoLys, Dpr(beta-Ala), alpha-(N-amidino-4'-piperidine)Gly, (4'-guanidino)Gly, (4'-guanidino)Phe, D-(4'-guanidino)Phe, beta-(N-amidino-4'-peperidine)Ala or homo-Ala-4'-pip(N-amidino); and

S₅ is Trp, Trp-OH, Trp-NH₂, Trp-Cys-NH₂, D-Trp, D-Trp-NH₂, Trp-Val-NH₂, 3'-Pya-NH₂, Phe-NH₂, MeTrp-NH₂, beta-Ala-Trp-NH₂, aminobutylamide, Nal 1-NH₂, D-Nal 1-NH₂, Nal 2-NH₂, D-

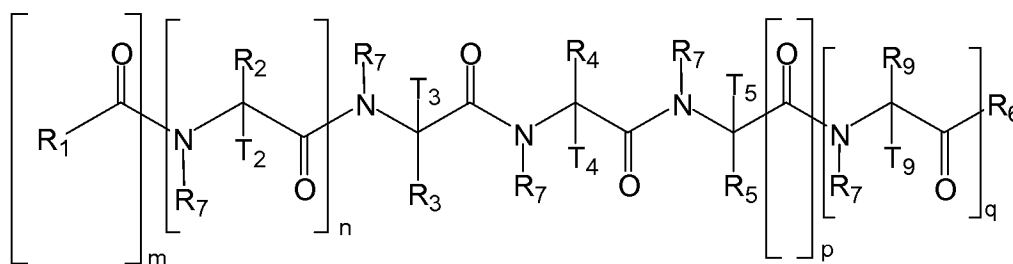
Nal 2-NH₂, Tic-NH₂, D-Tic-NH₂, 1'-aminoindan, 1'-aminoindane-1-carboxyl-NH₂, Aic-NH₂, Atc-NH₂, Disc-NH₂, Tpi-NH₂, D-Tpi-NH₂, Tiq-NH₂, D-Tiq-NH₂, tryptamide, NMe-tryptamide, alpha-Me-tryptamide, 2'-(4"-methylphenyl)ethylamide, 3',4'-Cl₂)phenylmethylamide, 3'-phenylpropylamide, 2',4'-dichlorobenzylamide, 3'-(1H-imidazol)propylamide, 4-phenyl-piperidine-4-carbonamide, 3-phenyl-1-propylamide, 2,4-dichlorophenethylamide, S-(-)-1-(2-naphthyl)ethylamide, S-(-)-1-(1-naphthyl)ethylamide, 2'-methylbenzylamide, 4'-methylbenzylamide, 2',2'-diphenylethylamide, 1-(2-pyridyl)piperazine, N-benzylmethylamide, histamide, R-(+)-1-(2-Naphthyl)ethylamide, Trp-Asp-NH₂, Trp-Asp-Phe-NH₂, Asp-Trp-NH₂, Ala-Trp-NH₂, Trp-Ala-NH₂, phenethylamide or Trp-Asp-OH.

14. (Original) The peptide of claim 13 consisting of the sequence

7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Ala-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-Phe-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-Asp-NH₂,
 heptanoyl-Thr(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Nal 2-Arg-Trp-NH₂,
 7'-amino-heptanoyl-Ala-D-Nal 2-Arg-Trp-NH₂,
 Ser(Bzl)-D-Nal 2-Arg-Trp-NH₂,
 Ser(Bzl)-D-Nal 2-Arg-D-Trp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-MeTrp-NH₂,
 heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-MeTrp-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-NMe-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-alpha-Me-Tryptamide,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-S-(-)-1-(1-Naphthyl)ethylamide,

7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 1-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-D-Nal 2-NH₂,
 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Nal 2-NH₂,
 2'-naphthylacetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂,
 4'-phenylbutyryl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂,
 3',4'-dichlorophenyl-acetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂, or
 3'-CF₃phenyl-acetyl-Ala-D-Phe(4-Cl)-Arg-Trp-NH₂.

15. (Withdrawn-currently amended) A melanocortin receptor-specific linear peptide of the formula:



where:

R₁ is an aliphatic L- or D-amino acid, N-acylated L- or D-aliphatic amino acid or R₈;

R₈ is independently selected from the group consisting of linear or branched C₁ to C₁₇ alkyl, aryl, heteroaryl, alkene, alkenyl, or aralkyl chains;

R₂ and R₃ are each independently H, CH₃, or an aromatic substituent aryl or heteroaryl side chain of a natural or synthetic L- or D-amino acid containing at least one aromatic ring moiety, wherein the ring(s) may additionally be functionalized by one or more halogen, alkyl or aryl groups;

R₄ is a positively charged aliphatic or aromatic side chain of a natural or synthetic L- or D-amino acid, wherein the chain comprises at least one nitrogen-containing group, or is a neutral aliphatic side chain having hydrogen donors and/or acceptors;

R₅ is H, CH₃, an aromatic substituent aryl or heteroaryl side chain of a natural or synthetic

L- or D-amino acid containing at least one aromatic ring moiety wherein the ring(s) may additionally be functionalized by one or more halogen, alkyl or aryl groups, or a substituent alkyl or hydrogen bonding polar side chain of a natural or synthetic L- or D-amino acid wherein the side chain has a hydrogen donor or acceptor moiety;

R_6 is hydroxide, NH_2 , or $NH-R_8$;

R_7 is H, methyl, ethyl, propyl, butyl, or a higher linear or branched chain terminating in an amino group, benzyl, or aralkyl group;

R_9 is H or an amino acid side chain group;

m is ~~normally 1~~ 1 or 0, with the proviso that if m is 0 then m may be 0 in which case this functionality is not present and the N-terminal group is an amine;

n is ~~normally 1~~ 1 or 0, with the proviso that if n is 0 then n may be 0 in which case this amino acid is not present;

p is ~~normally 1~~ 1 or 0 with the proviso that when p is 0 the chain terminates with the combination of R_5 and T_5 and there is no q and no R_6 ;

q is ~~normally 1~~ 1 or 0 with the proviso that when q is 0 and p is 1 then the terminal group is R_6 ; and

T_2 , T_3 , T_4 , T_5 , and T_9 are each H, CH_3 , C_2H_5 or a benzyl group;

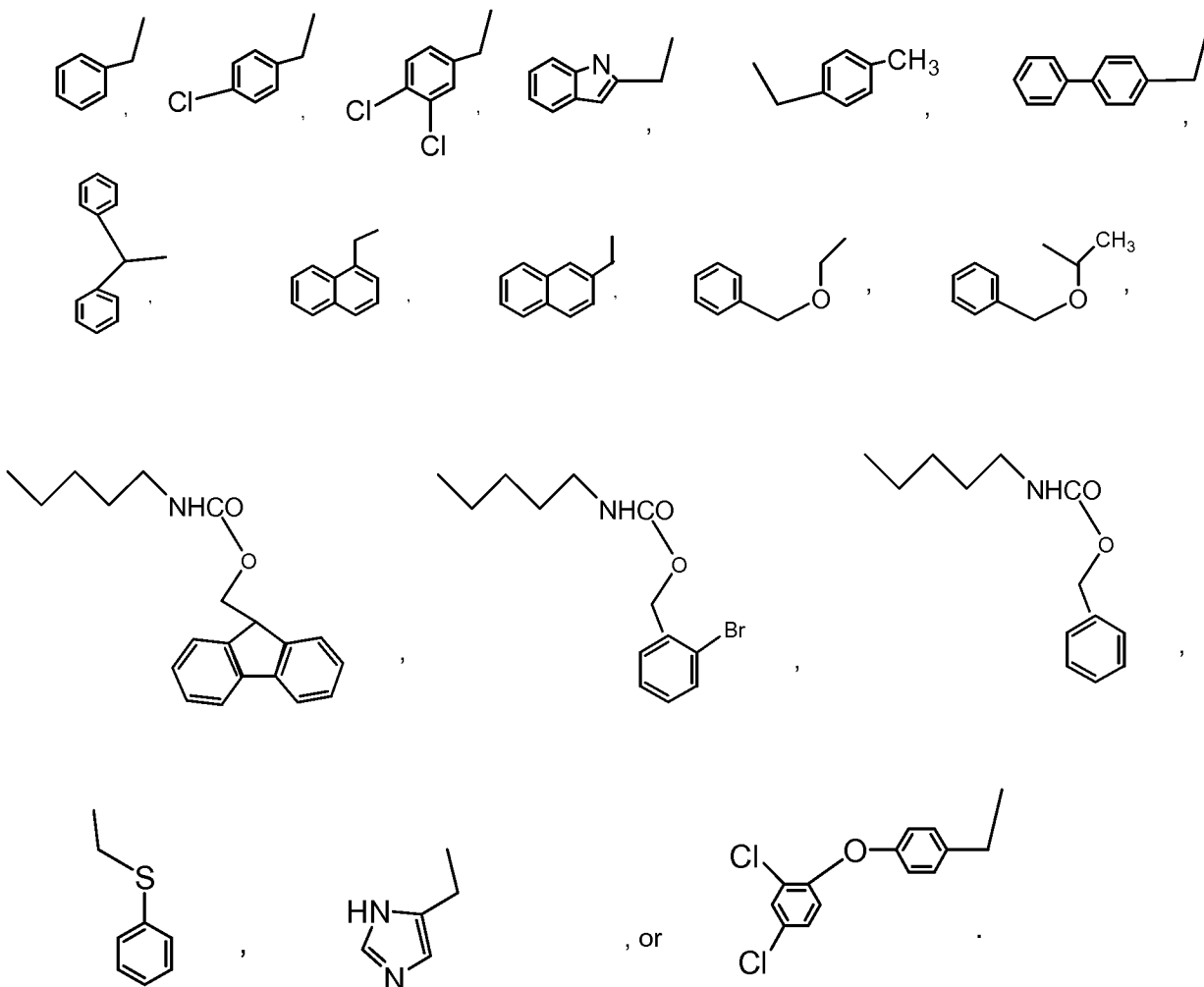
provided that one or more of the pairs R_2 and T_2 , or R_3 and T_3 , or R_4 and T_4 , or R_5 and T_5 , or R_9 and T_9 moieties may be joined together by additional carbon-carbon bonds to form a five-, six- or seven-membered ring structure; and

further provided that one or more of R_2 , R_4 , R_5 or R_9 may be joined to the R_7 group that immediately precedes such R_2 , R_4 , R_5 or R_9 group by additional carbon-carbon bonds to form a five-, six- or seven-membered ring structure, thereby fixing such R_2 , R_4 , R_5 or R_9 group to the immediately preceding nitrogen atom.

16. (Withdrawn) The linear peptide of claim 15 wherein R_8 is a C_1 to C_{17} aliphatic linear chain or branched chain group, an acylated group derived from C_1 to C_{17} aliphatic linear chain or branched chain group, an omega amino and carboxylic derivative of a C_1 to C_{17} aliphatic linear chain or branched chain groups, or an omega amino derivative of an acylated group

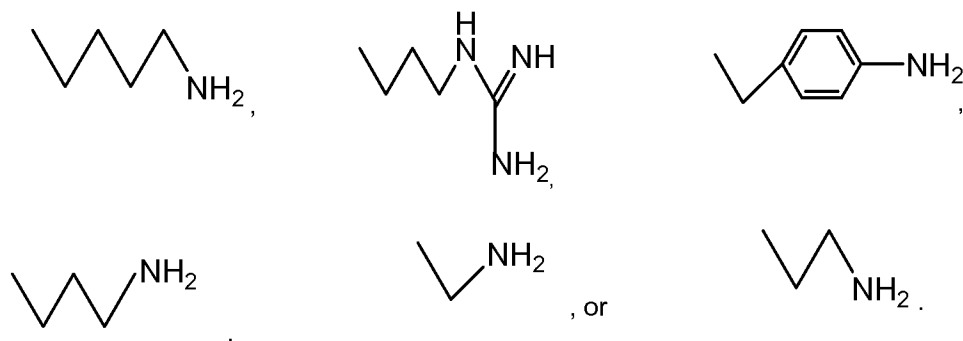
derived from a C₁ to C₁₇ aliphatic linear chain or branched chained group.

17. (Withdrawn) The linear peptide of claim 15 wherein at least one of R₂ and R₃ are

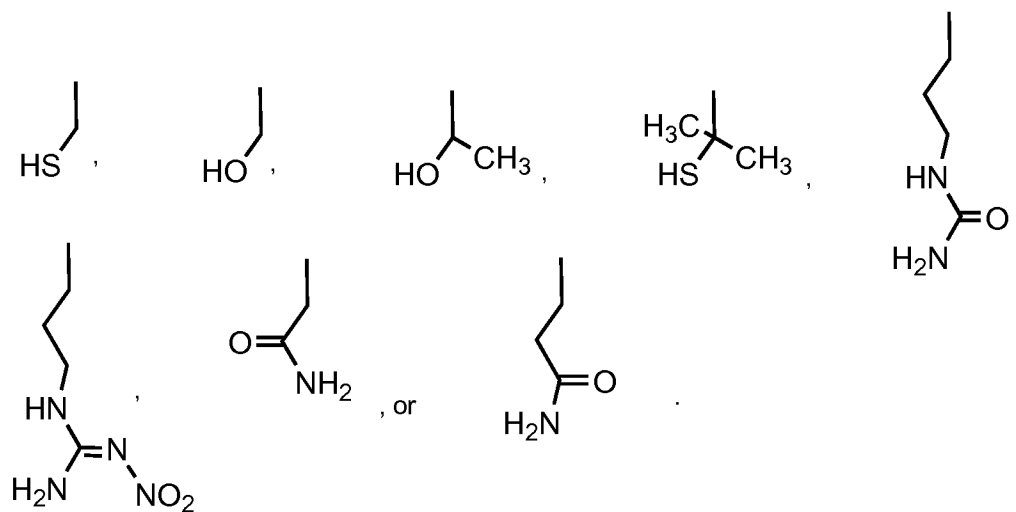


18. (Withdrawn) The peptide of claim 15 wherein R₄ is a positively charged aliphatic or aromatic side chain of a natural or synthetic L- or D-amino acid, wherein the at least one nitrogen-containing group in the chain is an amide, imide, amine or nitrile.

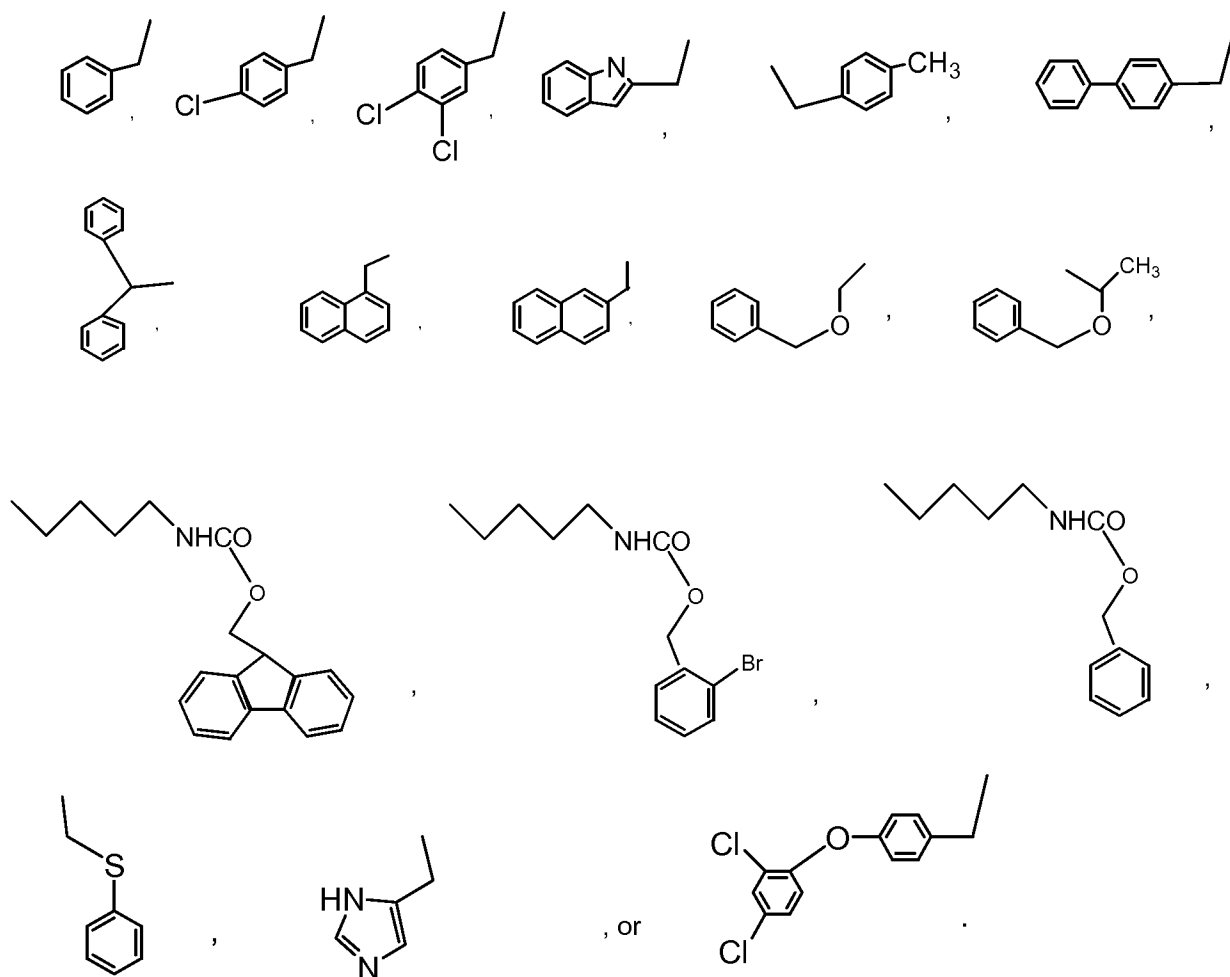
19. (Withdrawn) The peptide of claim 15 wherein R₄ is



20. (Withdrawn) The peptide of claim 15 wherein R₄ is a neutral aliphatic side chain having hydrogen donors and/or acceptors comprising:

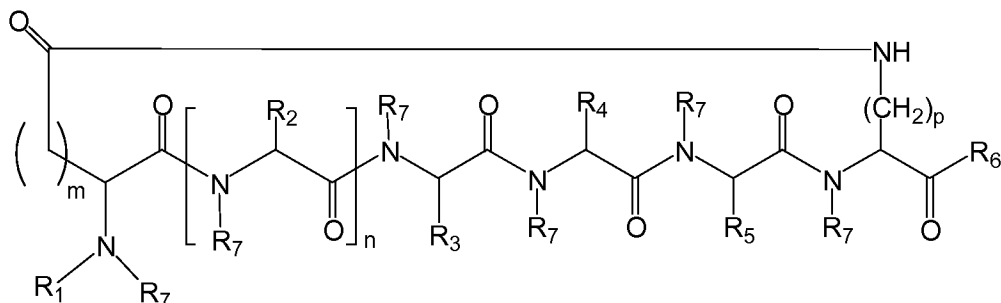


21. (Withdrawn) The peptide of claim 15 wherein R₅ is



22. (Withdrawn) The peptide of claim 15 wherein R₉ is methyl, ethyl, propyl, butyl, a higher linear or branched chain, or a linear chain terminating in an amino group, benzyl, or aralkyl group.

23. (Withdrawn-currently amended) A melanocortin receptor-specific cyclic peptide of the formula:



where:

R₁ is H, an aliphatic L- or D-amino acid, N-acylated aliphatic L- or D-amino acid or R₈;

R₂, R₃ and R₅ are independently each H, CH₃, an aromatic substituent aryl or heteroaryl side chain of a natural or synthetic L- or D-amino acid containing at least one aromatic moiety, wherein the ring(s) may additionally be functionalized by halogen, alkyl or aryl groups;

R₄ is a positively charged aliphatic or aromatic side chain of a natural or synthetic L- or D-amino acid, wherein the chain comprises at least one nitrogen-containing group, or is a neutral aliphatic side chain having hydrogen donors and/or acceptors;

R₆ is hydroxide, NH₂, or NH-R₈;

R₇ is H, methyl, ethyl, propyl, butyl, a higher linear or branched chain homolog, or a chain terminating in an amino group, benzyl, or aralkyl group;

R₈ is, in each instance, independently a linear or branched C₁ to C₁₇ alkyl, aryl, heteroaryl, alkene, alkenyl, or aralkyl chain;

m is 1 or 2;

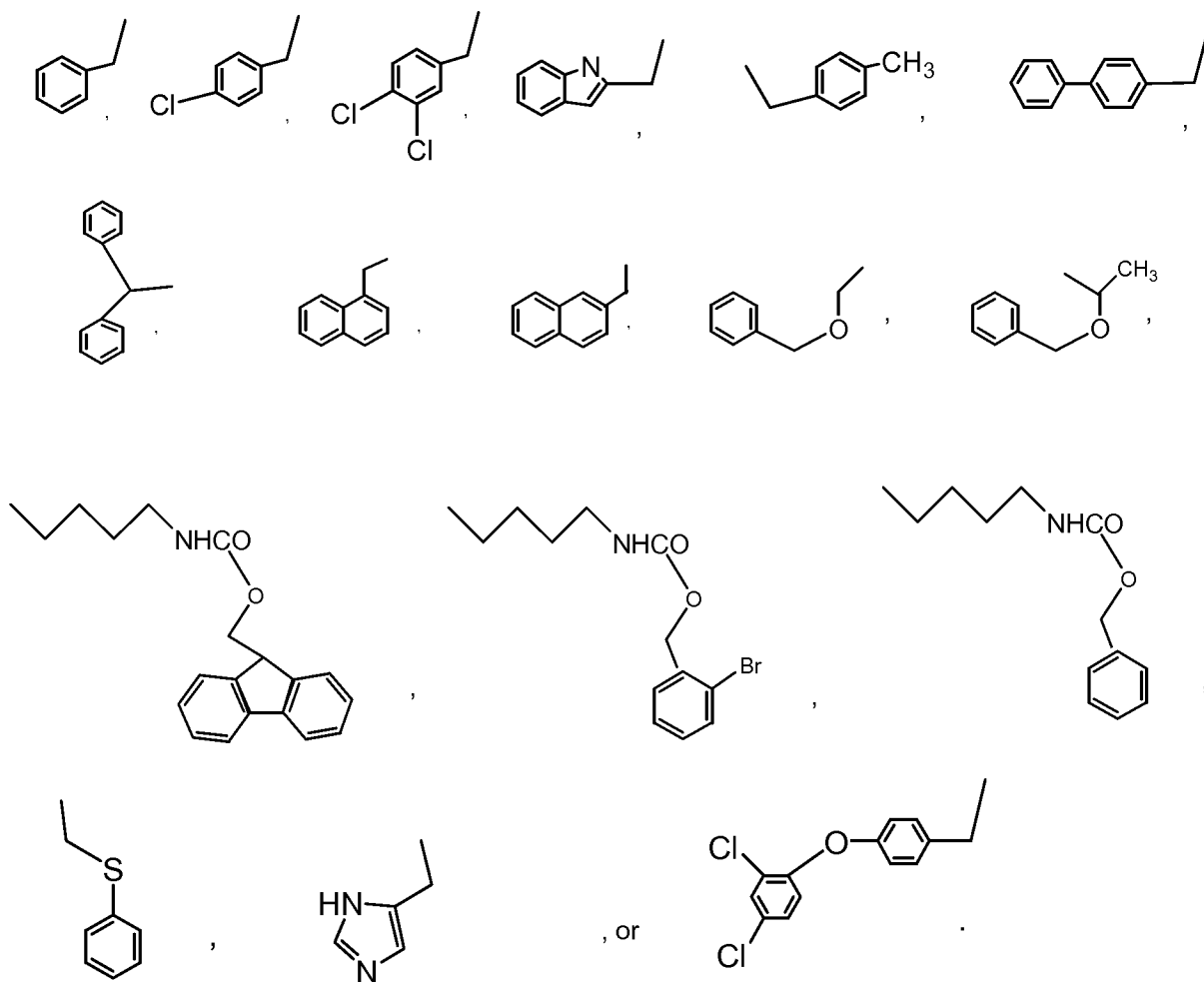
n is ~~normally 1~~ 1 or 0 with the proviso that if n is 0 then n may be 0 in which case this amino acid is not present; and

p is 1 to 5.

24. (Withdrawn) The cyclic peptide of claim 23 wherein R₈ is a C₁ to C₁₇ aliphatic

linear chain or branched chain group, an acylated group derived from C₁ to C₁₇ aliphatic linear chain or branched chain group, an omega amino and carboxylic derivative of a C₁ to C₁₇ aliphatic linear chain or branched chain groups, or an omega amino derivative of an acylated group derived from a C₁ to C₁₇ aliphatic linear chain or branched chain group.

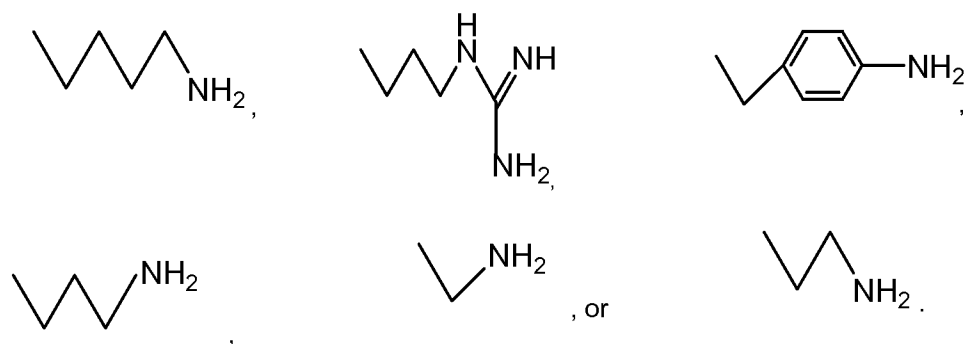
25. (Withdrawn) The cyclic peptide of claim 23 wherein at least one of R₂, R₃ or R₆ are independently



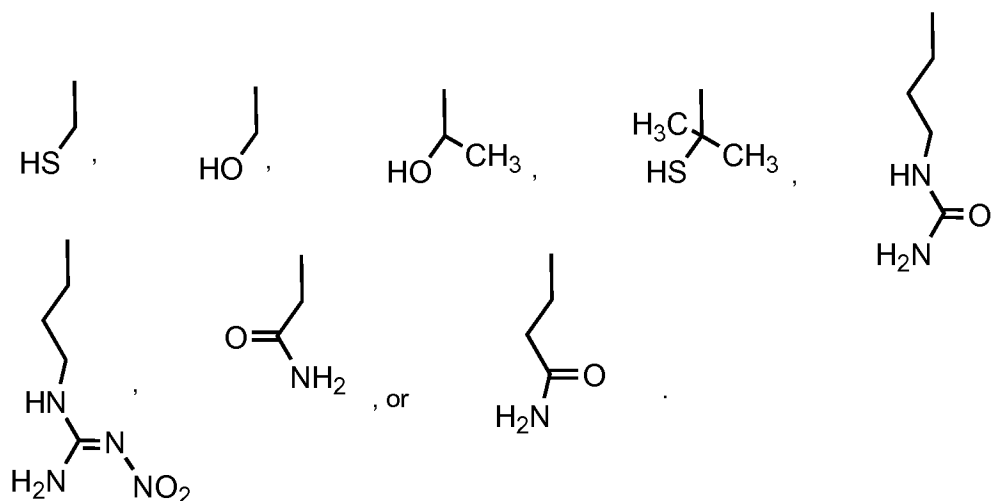
26. (Withdrawn) The cyclic peptide of claim 23 wherein R₄ is a positively charged aliphatic or aromatic side chain of a natural or synthetic L- or D-amino acid, wherein the at least

one nitrogen-containing group in the chain is an amide, imide, amine or nitrile.

27. (Withdrawn) The cyclic peptide of claim 23 wherein R₄ is



28. (Withdrawn) The cyclic peptide of claim 23 wherein R₄ is a neutral aliphatic side chain having hydrogen donors and/or acceptors comprising:



29. (Withdrawn) A method of stimulating sexual response in a mammal, comprising administering a pharmaceutically sufficient amount of a melanocortin receptor 3 and/or 4 selective agonist peptide of any of the foregoing claims.

30. (Withdrawn) The method of claim 29 comprising a method of administration selected from the group consisting of intravenous, subcutaneous, intramuscular, parenteral, intranasal, oral, dermal, inhalation, buccal, pulmonary, ocular, sublingual and vaginal administration.

31. (Withdrawn) A method of decreasing food intake in a mammal, comprising administering a pharmaceutically sufficient amount of a melanocortin receptor 4 and/or 5 selective agonist peptide of any of claims 1 to 28.

32. (Withdrawn) The method of claim 31 comprising a method of administration selected from the group consisting of intravenous, subcutaneous, intramuscular, parenteral, intranasal, oral, dermal, inhalation, buccal, pulmonary, ocular, sublingual and vaginal administration.

33. (Original) A peptide of the sequence 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂.

34. (Original) A pharmaceutical composition comprising a peptide of the sequence 7'-amino-heptanoyl-Ser(Bzl)-D-Phe(4-Cl)-Arg-Trp-NH₂.